 <b>NEONATAL CLINICAL PRACTICE GUIDELINES</b>	Title: <b>Management of Cardiorespiratory Events in Newborns</b>	
	Approval Date: June 2022	Pages: 1-8
	Approved by: Neonatal Patient Care Teams, HSC & SBGH Child Health Standards Committee	

## 1.0 PURPOSE AND INTENT

- 1.1 To provide a process for the management, monitoring, treatment, and documentation of cardiorespiratory events including apnea, bradycardia, and desaturations in premature infants in neonatal units who are not intubated.
- 1.2 To provide for safe, effective, and consistent use of caffeine to treat apnea of prematurity.

*Note: All recommendations are approximate guidelines only and practitioners must take in to account individual patient characteristics and situation. Concerns regarding appropriate treatment must be discussed with the attending neonatologist.*

## 2.0 PRACTICE OUTCOME

- 2.1 Prevent and minimize the potential long-term neurodevelopmental impact of these events.
- 2.2 Reduce length of stay and readmission rates.
- 2.3 Decrease interprovider variability.

## 3.0 DEFINITIONS

- 3.1 Clinically Significant Cardiopulmonary Event (CSCPE), include any one of the following:
  - a. Apneic event with a respiratory pause  $\geq 20$  seconds
  - b. Apneic event with a respiratory pause  $< 20$  seconds accompanied by a bradycardia  $< 80$  bpm for any duration or with a systemic oxygen desaturation  $< 85\%$  or accompanied by central cyanosis (if no pulse oximeter was being used) for any duration
  - c. Bradycardia: a heart rate  $< 80$  bpm for  $> 5$  seconds
  - d. Oxygen desaturation: systemic oxygen saturation (SpO<sub>2</sub>)  $< 80\%$  for  $> 5$  seconds or central cyanosis (if no pulse oximeter was being used).
- 3.2 Apnea of Prematurity: Apneas occurring in infants less than 37 weeks post-menstrual age (PMA) in the absence of any other condition that may cause apnea (i.e. sepsis).
- 3.3 Periodic Breathing: A series of 3 or more respiratory pauses per minute, each longer than 3 seconds with intervals of respiration between pauses. This pattern of breathing is common in newborns and is usually not associated with significant bradycardia or desaturation. In those with decreased pulmonary reserve periodic breathing may culminate in oxygen desaturation albeit brief.
- 3.4 Bradycardia:
  - 3.4.1 Infant current weight less than 1250 grams
    - 3.4.1.1 A heart rate less than 100 beats per minute
  - 3.4.2 Infant current weight greater than 1250 grams
    - 3.4.2.1 A heart rate less than 80 beats per minute in term infants

- 3.4.3 Profound Bradycardia
  - 3.4.3.1 Heart rate less than 60 beats per minute for 5 seconds or more irrespective of weight).
- 3.5 Desaturation: Oxygen saturation (SpO<sub>2</sub>) less than 85% for 10 seconds or more unless otherwise specified in physician's/NNP's order.
  - 3.5.1 Hypoxemia: Oxygen Saturation (SpO<sub>2</sub>) less than 80% for 10 seconds or more
  - 3.5.2 Intermittent hypoxemic events are desaturations below 80% for more than 10 seconds and/or an abnormal histogram in an infant in room air with more than 10% of the time below 90%).
- 3.6 Respiratory Support: An artificial means of mechanical support which includes intubation, tracheostomy, nCPAP (nasal continuous positive airway pressure) or non-invasive ventilation, an oral or nasal airway or low flow oxygen (i.e., nasal prongs) (regardless of oxygen level).
- 3.7 Positive Pressure Ventilation (PPV): the application of positive pressure using a self-inflating or flow inflating anaesthesia bag through a full-face mask encompassing the nose and mouth.
- 3.8 Stimulation is of four types as follows:
  - a. Self: no stimulation was required by a caregiver for the infant to recover from the event.
  - b. Gentle: gentle tactile stimulation such as touching or rubbing of the back was required by a caregiver for the infant to recover from the event.
  - c. Moderate: flicking the sole of the foot, re-positioning the infant side to side, using blow-by oxygen was required by the caregiver for the infant to recover from the event.
  - d. Vigorous: ventilation with bag and mask or greater resuscitative efforts were required by a caregiver for the infant to recover from the event
- 3.9 Self-Resolved Event: Infant recovered spontaneously with respect to respiration and oxygen saturation.
- 3.10 Mild Event: Infant recovered respiration and oxygenation recovery within 10 seconds of minimal tactile stimulation.
- 3.11 Moderate Event: Infant recovered respiration and oxygenation within 30 seconds of the beginning of the event after moderate tactile stimulation.
- 3.12 Severe Event: Infant did not show recovery of respiration and oxygenation within 30 seconds of the beginning of the event in spite of moderate tactile stimulation.
- 3.13 Gestational Age (GA) (completed weeks, not rounded up): Time elapsed between the first day of the last menstrual period and the day of delivery. If pregnancy was achieved using assisted reproductive technology, gestational age is calculated by adding 2 weeks to the conceptual age.
- 3.14 Chronological Age (days, weeks, months, or years): Time elapsed since birth.
- 3.15 Postmenstrual Age (PMA): Gestational age plus chronological age. This is the preferred term to use during the perinatal neonatal hospital stay.

- 3.16** Preterm: A preterm (premature) infant is one born at a gestational age of 259 days (37 weeks) or less.
- 3.17** Methylxanthines: Methylated derivatives of xanthine, including caffeine, theobromine and theophylline that serve as a smooth muscle relaxant and cardiac muscle and central nervous system stimulant.

## **4.0** **GUIDELINES**

### **4.1** **General Management**

Provide cardiorespiratory monitoring, including oxygen saturation histograms, to all infants who are less than 35 weeks 0 days gestational age. Cardiorespiratory monitoring, including oxygen saturation histograms, should be continued for at least 72 hours from the last documented cardiorespiratory episode. If the infant was on caffeine, continue cardiorespiratory monitoring, including daily histograms for at least 7 days after discontinuation of caffeine.

If the infant has frequent cardiorespiratory episodes or significant increase in respiratory support, or failure to wean the present respiratory support, a full clinical examination should be considered including assessment of airway patency, thermal stability, blood pressure, glucose regulation and Lung Ultrasound (LUS)/Chest XR. Other potential causes of apnea should also be considered.

Consider treatment with methylxanthine medications (see section below) or CPAP.

Consider a control of breathing study in infants meeting all the following inclusion criteria:

- a. Having persistent apneas, bradycardias, and/or desaturations after a corrected post conceptional age greater than 36 weeks 6 days.
- b. Feeding ad lib demand.
- c. Not on caffeine maintenance.
- d. No other medical condition keeping the infant from being discharged home.

If an immature breathing pattern is recognized during the sleep study that improves after a loading dose of caffeine, consider discharge home on caffeine after a few days of observation in the unit at the discretion of the neonatologist.

### **4.2** **Management of Events**

When the monitor alarms, evaluate the infant for airway patency, noting if hyperextension or flexion of the neck is present. Reposition the head and neck if necessary. Maintain the head and neck in a neutral position. Additionally, respiratory movements, color, and heart rate should be assessed.

- 4.2.1** If the infant continues to be apneic or bradycardic after the assessment and position change, initiate appropriate tactile stimulation.
- 4.2.2** If the infant continues to have inadequate respiratory effort after 30 seconds from the beginning of the event, discontinue stimulation, administer positive pressure ventilation (PPV) **and call for assistance**. If the infant has some respiratory effort, provide only Positive End Expiratory Pressure (PEEP) of 5-6 cmH<sub>2</sub>O. If the infant shows no respiratory effort, provide breaths at a rate of 40-60 per minute. Avoid hyperventilation as this may decrease the infant's pCO<sub>2</sub> level and suppress the stimulus to breathe triggering apnea itself. Administering free-flow oxygen or continuing to provide tactile stimulation to an apneic or

bradycardic infant will not treat the resultant hypoxemia as no oxygen is being delivered to the alveoli.

- 4.2.3 Reassess respiratory effort, heart rate and oxygen saturation ongoing once PPV is established. Increase the inspired oxygen (FiO<sub>2</sub>) only if the baby's heart rate and oxygen saturation do not show any improvement at this point.
- 4.2.4 Continue PPV until regular respiration and normal heart rate have returned to baseline.
- 4.2.5 Document on the Neonatal Cardiorespiratory Events Record:
  - a. Any respiratory pause that is greater than 20 seconds or shorter periods with an associated bradycardia or desaturation not caused by care/interventions (i.e., suctioning, or gastric tube insertion).
  - b. The duration of the episode from the time that the monitor indicates the episode began (using the events record or Oxygen feature) until the episode resolves.
- 4.2.6 DO NOT RECORD on the Neonatal Cardiorespiratory Events Record:
  - a. Episodes of periodic breathing, bradycardia, or respiratory pauses less than 20 second duration that are not associated with significant bradycardia or desaturation.
  - b. Decreased oxygen saturations that are not associated with a respiratory pause or bradycardia.
  - c. Transient episodes of bradycardia that are self-resolved within 10 seconds and are not associated with a desaturation below 85% or respiratory pause.
  - d. Any events that were not witnessed, assessed, and responded to, but were found only on the monitor on retrospective review.
  - e. Any events caused by interventions such as gastric tube insertion or suctioning.
  - f. Any events reflective of monitor artifact.
  - g. Such events as described in sections a-f while not recorded may be discussed on patient rounds as to whether they require further investigation or not.

### 4.3 Caffeine for Apnea of Prematurity

- 4.3.1 Intravenous (IV) and oral Caffeine is ordered as 'Caffeine', not caffeine citrate.

Adjust dosing and frequency according to the following:

- a. Oral and IV doses are the same.
  - b. Usual loading dose: 10 mg/kg.
  - c. If there is an ineffective clinical response to the initial load there may be additional loading doses at the discretion of the Neonatologist.
  - d. Usual maintenance dose: 2.5-5 mg/kg/day given once a day.
  - e. A higher maintenance dose may be considered, with the neonatologist approval, especially in extreme low gestational age infants to prevent reintubation
- 4.3.2 Additional loading doses of 5 mg/kg may be given after 4 hours up to maximum of 20 mg/kg/day total.

- 4.3.3 Consider discontinuing caffeine between 32- and 34-weeks PMA if: a) the infant is off NCPAP b) no significant or frequent events are observed; and c) daily histogram is considered acceptable by the team.
- 4.3.4 Continue cardiorespiratory monitoring for 5 to 7 days after discontinuing caffeine and until the baby demonstrates no episodes of apnea for at least 3 days. Note: Caffeine half life has a mean of approximately 100 hours in the preterm infant with longer half lives at the lowest gestational ages at birth. (REF: Abdel-Hady H, Nasef N, Shabaan AE, Nour I. Caffeine therapy in preterm infants. *World journal of clinical pediatrics*. 2015;4(4):81-93.

## 5.0 Discharge criterion

- 5.1 Off caffeine for 5-7 days and no CSCPE.
- 5.2 72 hours CSCPE free post last CSCPE.
- 5.3 For those who have met all goals for discharge but continue to demonstrate a need for caffeine therapy, discharge may be done with planned continuation of caffeine in the home. If this is being considered, consult with the neonatal control of breathing lab prior to discharge.

## 6.0 REFERENCES

- 6.1 American Heart Association (2017). *Textbook of Neonatal Resuscitation 7th Edition*.
- 6.2 Barbara Schmidt, Robin S Roberts, Peter J Anderson, Elizabeth V Asztalos, Lorrie Costantini, Peter G Davis, Deborah Dewey, Judy D'Ilario, Lex W Doyle, Ruth E Grunau, Diane Moddemann, Harvey Nelson, Arne Ohlsson, Alfonso Solimano, Win Tin, Caffeine for Apnea of Prematurity (CAP) Trial Group, Academic Performance, Motor Function, and Behavior 11 Years After Neonatal Caffeine Citrate Therapy for Apnea of Prematurity: An 11-Year Follow-up of the CAP Randomized Clinical Trial, PMID: 28437520 DOI: 10.1001/jamapediatrics.2017.0238
- 6.3 Carole L Marcus, Lisa J Meltzer, Robin S Roberts, Joel Traylor, Joanne Dix, Judy D'Ilario, Elizabeth Asztalos, Gillian Opie, Lex W Doyle, Sarah N Biggs, Gillian M Nixon, Indra Narang, Rakesh Bhattacharjee, Margot Davey, Rosemary S C Horne, Maureen Cheshire, Jeremy Gibbons, Lorrie Costantini, Ruth Bradford, Barbara Schmidt, Caffeine for Apnea of Prematurity–Sleep Study, Long-term effects of caffeine therapy for apnea of prematurity on sleep at school age, PMID: 25171195 PMID: PMC4299611 DOI: 10.1164/rccm.201406-1092OC
- 6.4 Chen Kou, Dong Han, Zhaona Li, Wei Wu, Zunjie Liu, Yanan Zhang, Zhengping Gao, Influence of prevention of caffeine citrate on cytokine profile and bronchopulmonary dysplasia in preterm infants with apnea, April 2020, PMID: 30961342 DOI: 10.23736/S0026-4946.19.05428-8
- 6.5 Davis, P.G., Schmidt, B., Roberts, R.S., Doyle, L.W., Asztalos, E., Haslam, R., Sinha, S., tin, W. & Caffeine for Apnea of Prematurity Trial Group. (2010) Caffeine for apnea of prematurity trial: benefits may vary in subgroups. *Journal of Pediatrics*, 156(3), 382-387.
- 6.6 Eilan Alhersh, Dina Abushanab, Samaher Al-Shaibi, Daoud Al-Badriyeh, Caffeine for the Treatment of Apnea in the Neonatal Intensive Care Unit: A Systematic Overview of Meta-Analyses, August 2020, PMID: 32488731 PMID: PMC7266675 DOI: 10.1007/s40272-020-00404-4
- 6.7 Gillies, D., & Wells, D. (2010). Positioning for acute respiratory distress in hospitalized infant and children, *Cochrane Database*, 2.

- 6.8** Gray, P.H., Flenady, V.J., Charles, B.G., Steer, P.A. & Caffeine Collaborative Study Group (2011). Caffeine citrate for very preterm infants. Effects on development, temperament, and behavior. *Journal of Pediatrics and Child Health*. 47(4), 167-172.
- 6.9** Henderson-Smart, D.J. & De Paoli, A.G. (2010). Methylxanthine treatment for apnoea in preterm infants. *Cochrane Database*. Dec 8(12), CD000140.
- 6.10** Henderson-Smart, D.J. & De Paoli, A.G. (2010). Prophylactic methylxanthine for prevention of apnoea in preterm infants. *Cochrane Database*. Dec 8(12), CD000432.
- 6.11** Hodgman, J. E., Gonzalez, F., Hoppenbrouwers, T., & Cabal, L. A. (1990). Apnea, transient episodes of bradycardia, and periodic breathing in preterm infants. *American Journal of Diseases in Children*, 144(1), 54-57.
- 6.12** Hye Won Park, Gina Lim, Sung-Hoon Chung, Sochung Chung, Kyo Sun Kim, Soo-Nyung Kim, Early Caffeine Use in Very Low Birth Weight Infants and Neonatal Outcomes: A Systematic Review and Meta-Analysis, November 2015, PMID: 26713059 PMCID: PMC4689828 DOI: 10.3346/jkms.2015.30.12.1828
- 6.13** Ines M. Mürner-Lavanchy, Lex W. Doyle, Barbara Schmidt, Robin S. Roberts, Elizabeth V. Asztalos, Lorrie Costantini, Peter G. Davis, Deborah Dewey, Judy D'Ilario, Ruth E. Grunau, Diane Moddemann, Harvey Nelson, Arne Ohlsson, Alfonso Solimano, Win Tin, Peter J. Anderson and for the Caffeine for Apnea of Prematurity (CAP) Trial Group, Neurobehavioral Outcomes 11 Years After Neonatal Caffeine Therapy for Apnea of Prematurity. Double-blind, randomized, placebo-controlled trial. *Pediatrics* May 2018, 141 (5) e20174047; DOI: <https://doi.org/10.1542/peds.2017-4047>
- 6.14** Jing Chen , Lu Jin , Xiao Chen, Efficacy and Safety of Different Maintenance Doses of Caffeine Citrate for Treatment of Apnea in Premature Infants: A Systematic Review and Meta-Analysis, PMID: 30671477 PMCID: PMC6323495 DOI: 10.1155/2018/9061234
- 6.15** K Kreutzer, D Bassler, Caffeine for apnea of prematurity: a neonatal success story, PMID: 24931325 DOI: 10.1159/000360647
- 6.16** Keene, D.J., Wimmer, J.E. Jr. & Mathew, O.P., (2000). Does supine positioning increase apnea, bradycardia, and desaturation in preterm infants? *Journal of Perinatology*, 20(1), 17-20.
- 6.17** Laura Moschino, Sanja Zivanovic, Caroline Hartley, Daniele Trevisanuto, Eugenio Baraldi, Charles Christoph Roehr, Caffeine in preterm infants: where are we in 2020?, March 2020, PMID: 32154294 PMCID: PMC7049734 DOI: 10.1183/23120541.00330-2019
- 6.18** Lister, G. Rybin, D., Colton, T., Heeren, T., Hunt, C., Colson, E., Willinger, M., & Corwin, M. (2012). Relationship between Sleep Position and Risk of Extreme Cardiorespiratory Events. *The Journal of Pediatrics*.161(1). 22-25.
- 6.19** Lizhong Du, Xiaomei Tong, Chao Chen, Xirong Gao, Alessandra Gagnatelli, Jingyang Li, Debora Santoro, Sara Nicolardi, Laura Fabbri, Peyona, Caffeine Citrate for Apnea of Prematurity: A Prospective, Open-Label, Single-Arm Study in Chinese Neonates, March 2020, PMID: 32219085 PMCID: PMC7078308 DOI: 10.3389/fped.2020.00076
- 6.20** M Shivakumar, P Jayashree, Muhammad Najih, Leslie Edward Simon Lewis, Ramesh Bhat Y, Asha Kamath, - Shashikala, Comparative Efficacy and Safety of Caffeine and Aminophylline for Apnea of Prematurity in Preterm ( $\leq 34$  weeks) Neonates: A Randomized Controlled Trial, April 2017, PMID: 28474588 DOI: 10.1007/s13312-017-1088-0
- 6.21** Martin, R. J., Abu-Shaweesh, J. M., & Baird, T. M. (2004). Apnoea of prematurity. *Pediatric Respiratory Reviews*, (5), S377-S382.

- 6.22** P Chandrasekharan, M Rawat, A M Reynolds, K Phillips & S Lakshminrusimha, Apnea, bradycardia and desaturation spells in premature infants: impact of a protocol for the duration of 'spell-free' observation on interprovider variability and readmission rates, PMID: 29120450 PMCID: PMC5775039 DOI: 10.1038/jp.2017.174
- 6.23** Petter Brattström, Chiara Russo, David Ley, Matteo Bruschetti, High-versus low-dose caffeine in preterm infants: a systematic review and meta-analysis, March 2019, PMID: 30242903 DOI: 10.1111/apa.14586
- 6.24** Pichardo, R., Adam, J.S., Rosow, E., Bronzino, J. & Eisenfeld, L. (2003). Vibro-tactile stimulation system to treat apnea of prematurity. *Biomedical Instrument Technology*, 37(1), 34-40.
- 6.25** Pichler, G., Schmolzer, G., Muller, W. & Urlesberger, B. (2001). Body position-dependent changes in cerebral hemodynamics during apnea in preterm infants. *Brain Development*, 23(6), 395-400.
- 6.26** Rami A Ballout, Jann P Foster, Lara A Kahale, Lina Badr, Body positioning for spontaneously breathing preterm infants with apnoea, Review Cochrane Database Systematic Review 2017 Jan 9;1(1):CD004951.doi:10.1002/14651858.CD004951.pub3.
- 6.27** Roos Vliegthart, Martijn Miedema, Gerard J Hutten, Anton H van Kaam, Wes Onland, High versus standard dose caffeine for apnoea: a systematic review, November 2018, PMID: 29437799 DOI: 10.1136/archdischild-2017-313556
- 6.28** Sameh Mohammed, Islam Nour, Abd Elazeez Shabaan, Basma Shouman, Hesham Abdel-Hady, Nehad Nasef, High versus low-dose caffeine for apnea of prematurity: a randomized controlled trial, PMID: 25644724 DOI: 10.1007/s00431-015-2494-8
- 6.29** Sanjiv B Amin, Erica Burnell, Monitoring apnea of prematurity: validity of nursing documentation and bedside cardiorespiratory monitor, PMID: 23254381 PMCID: PMC4285412 DOI: 10.1055/s-0032-1329694
- 6.30** Schmidt, B, Anderson, P.J., Doyle, L.W., Dewey, D., Grunau, R.E., Asztalos, E.V., Davis, P.G., Tin, W., Moddemann, D., Solimano, A., Ohlsson, A., Barrington, K.J. & Roberts, R.S. (2012). Survival without disability to age 5 years after neonatal caffeine therapy for apnea of prematurity. *JAMA*, 307(3), 275-282.
- 6.31** Schoen, K., Yu, T., Stockman, C., Spigarelli, M.G. & Sherwin, C.M. (2014) Use of methylxanthine therapies for the treatment and prevention of apnea of prematurity. *Paediatric Drugs*. 16(2), 169-177
- 6.32** Spitzer, A.R. (2012). Evidence-based methylxanthine use in the NICU. *Clinics in Perinatology*. 39(1), 137-148.
- 6.33** Whyte, R. K. (2012). Neonatal management and safe discharge of late and moderate preterm infants. *Seminars in Fetal and Neonatal Medicine*, (17), 153-158.
- 6.34** Xiao Zhang, Hai-Tao Zhang, Yong Lyu, Li-Feng Wang, Zhen-Ying Yang, Clinical effect and safety of different maintenance doses of caffeine citrate in treatment of apnea in very low birth weight preterm infants: a prospective randomized controlled trial, PMID: 31208509 PMCID: PMC7389581 DOI: 10.7499/j.issn.1008-8830.2019.06.011
- 6.35** Ying Zhao, Xiuying Tian, Ge Liu, Clinical effectiveness of different doses of caffeine for primary apnea in preterm infants, PMID: 26791921 DOI: 10.3760/cma.j.issn.0578-1310.2016.01.008
- 6.36** Zhao, J., Gonzalez, F., & Dezhi, M. (2011). Apnea of prematurity: From cause to treatment. *European Journal of Pediatrics*, 170, 1097-1105.

**7.0 PRIMARY AUTHORS**

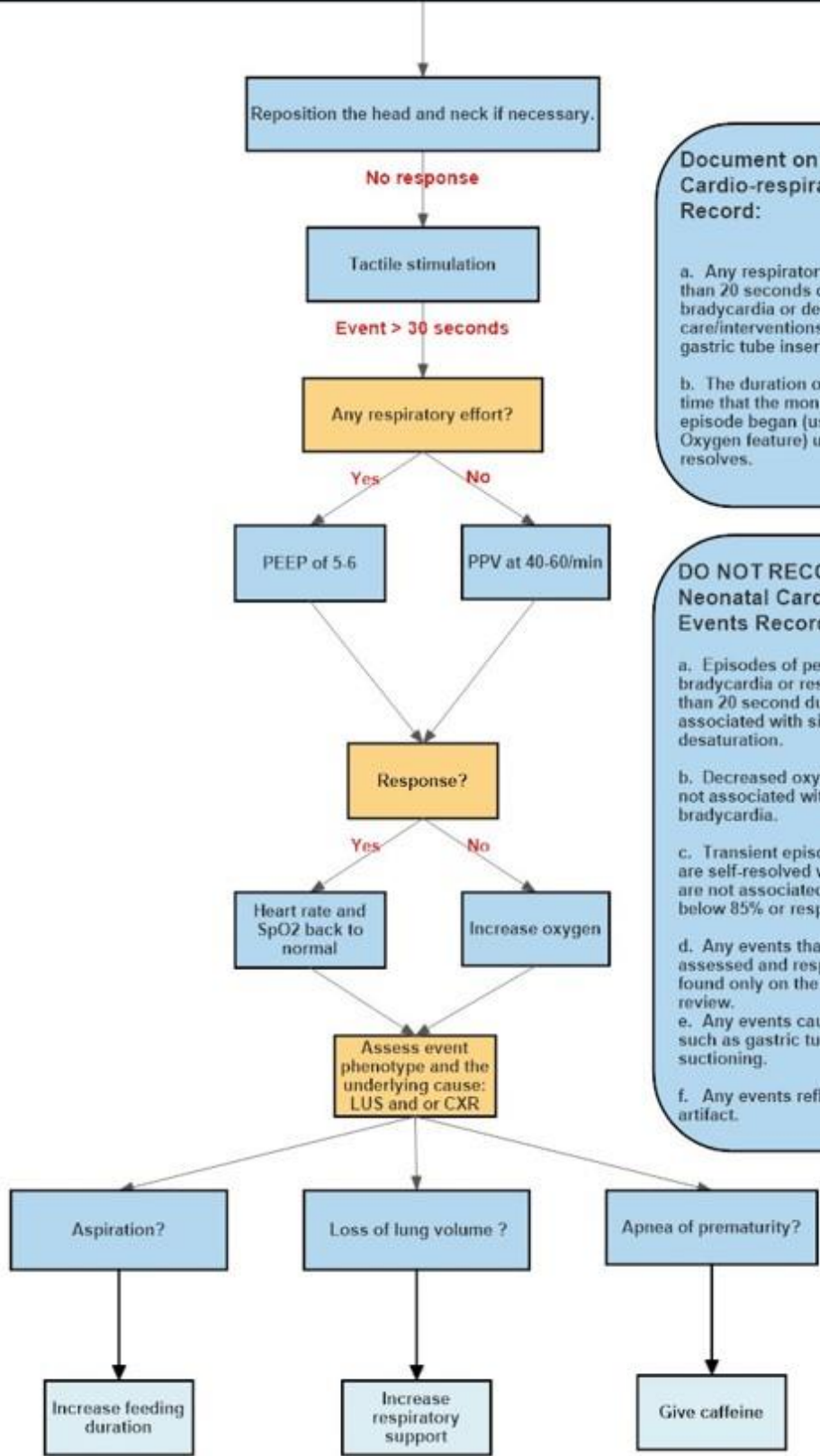
- 7.1 Dr. Asem Lashin Neonatology/Perinatology fellow PGY6
- 7.2 Dr. Ruben Alvaro, Assistant Medical Director NICU St. Boniface Hospital
- 7.3 Dr. Yasser El Sayed, Hemodynamics and POCUNEO Director, NICU, Health Sciences Centre
- 7.4 Dr. Michael Narvey, Section Head of Neonatology
- 7.5 Nicole Kjartanson, Manager of Patient Care / Education Instructor, NICU, Health Sciences Centre
- 7.6 Tanya Tichon, Nurse Educator, Health Sciences Centre
- 7.7 Natasha Macsych, Nurse Educator, Health Sciences Centre



**APPENDIX A – CARDIO-RESPIRATORY EVENT MANAGEMENT ALGORITHM**

**ALARM**

- Apneic event with a respiratory pause >20 seconds.
- Bradycardia < 80 BPM for > 10 seconds.
- Oxygen desaturation: oxygen saturation (SpO<sub>2</sub>) < 80% for > 5 seconds or central cyanosis.



**Document on the Neonatal Cardio-respiratory Events Record:**

- a. Any respiratory pause that is greater than 20 seconds or has an associated bradycardia or desaturation not caused by care/interventions (i.e., suctioning or gastric tube insertion).
- b. The duration of the episode from the time that the monitor indicates the episode began (using the events record or Oxygen feature) until the episode resolves.

**DO NOT RECORD on the Neonatal Cardio-respiratory Events Record:**

- a. Episodes of periodic breathing, bradycardia or respiratory pauses less than 20 second duration that are not associated with significant bradycardia or desaturation.
- b. Decreased oxygen saturations that are not associated with a respiratory pause or bradycardia.
- c. Transient episodes of bradycardia that are self-resolved within 10 seconds and are not associated with a desaturation below 85% or respiratory pause.
- d. Any events that were not witnessed, assessed and responded to, but were found only on the monitor on retrospective review.
- e. Any events caused by interventions such as gastric tube insertion or suctioning.
- f. Any events reflective of monitor artifact.